

**AMPICILLIN AND SULBACTAM - ampicillin sodium and sulbactam sodium injection, powder, for solution**  
Baxter Healthcare Corporation

**PHARMACY BULK PACKAGE NOT FOR DIRECT INFUSION**

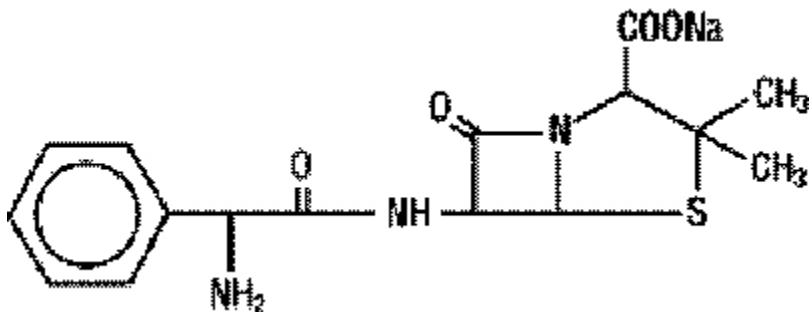
**Rx only**

To reduce the development of drug-resistant bacteria and maintain the effectiveness of Ampicillin and Sulbactam for Injection and other antibacterial drugs, Ampicillin and Sulbactam for Injection should be used only to treat or prevent infections that are proven or strongly suspected to be caused by bacteria.

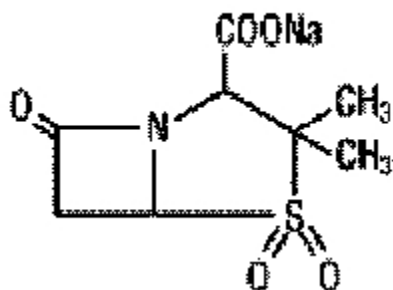
**DESCRIPTION**

Ampicillin and Sulbactam for Injection, USP is an injectable antibacterial combination consisting of the semisynthetic antibiotic ampicillin sodium and the beta-lactamase inhibitor sulbactam sodium for intravenous and intramuscular administration. **HOWEVER, THE INTENT OF THE PHARMACY BULK PACKAGE IS FOR PREPARATION OF SOLUTIONS FOR IV INFUSION ONLY.**

Ampicillin sodium is derived from the penicillin nucleus, 6-aminopenicillanic acid. Chemically, it is Monosodium D-(–)-6-(2-amino-2-phenylacetamido)-3,3-dimethyl-7-oxo-4-thia-1-azabicyclo[3.2.0]heptane-2-carboxylate and has a molecular weight of 371.39. Its chemical formula is  $C_{16}H_{18}N_3NaO_4S$ . The structural formula is:



Sulbactam sodium is a derivative of the basic penicillin nucleus. Chemically, sulbactam sodium is sodium penicillinate sulfone; sodium (2*S*, 5*R*)-3,3-dimethyl-7-oxo-4-thia-1-azabicyclo[3.2.0]heptane-2-carboxylate 4,4-dioxide. Its chemical formula is  $C_8H_{10}NNaO_5S$  with a molecular weight of 255.22. The structural formula is:



Ampicillin and Sulbactam for Injection is available as a white to off-white dry powder for reconstitution. Ampicillin and Sulbactam for Injection dry powder is freely soluble in aqueous diluents to yield pale yellow to yellow solutions containing ampicillin sodium and sulbactam sodium equivalent to 250 mg ampicillin per mL and 125 mg sulbactam per mL. The pH of the solutions is between 8.0 and 10.0.

Dilute solutions (up to 30 mg ampicillin and 15 mg sulbactam per mL) are essentially colorless to pale yellow. The pH of dilute solutions remains the same.

Each sterile Pharmacy Bulk Package contains 15 g Ampicillin and Sulbactam for Injection (10 g ampicillin as the sodium salt plus 5 g sulbactam as the sodium salt) and approximately 1150 mg (50 mEq) sodium. Ampicillin and Sulbactam for Injection Pharmacy Bulk Package is a vial containing a sterile preparation of ampicillin sodium and sulbactam sodium for parenteral use that contains many single doses. The Pharmacy Bulk Package is for use in a pharmacy admixture setting; it provides many single doses of ampicillin/sulbactam for addition to suitable parenteral fluids in the preparation of admixtures for intravenous infusion. (See DIRECTIONS FOR PROPER USE OF PHARMACY BULK PACKAGE.) FURTHER DILUTION IS REQUIRED BEFORE USE.

## CLINICAL PHARMACOLOGY

### General

Immediately after completion of a 15-minute intravenous infusion of Ampicillin and Sulbactam for Injection, peak serum concentrations of ampicillin and sulbactam are attained. Ampicillin serum levels are similar to those produced by the administration of equivalent amounts of ampicillin alone. Peak ampicillin serum levels ranging from 109 to 150 mcg/mL are attained after administration of 2000 mg of ampicillin plus 1000 mg sulbactam and 40 to 71 mcg/mL after administration of 1000 mg ampicillin plus 500 mg sulbactam. The corresponding mean peak serum levels for sulbactam range from 48 to 88 mcg/mL and 21 to 40 mcg/mL, respectively. After an intramuscular injection of 1000 mg ampicillin plus 500 mg sulbactam, peak ampicillin serum levels ranging from 8 to 37 mcg/mL and peak sulbactam serum levels ranging from 6 to 24 mcg/mL are attained.

The mean serum half-life of both drugs is approximately 1 hour in healthy volunteers.

Approximately 75 to 85% of both ampicillin and sulbactam are excreted unchanged in the urine during the first 8 hours after administration of Ampicillin and Sulbactam for Injection to individuals with normal renal function. Somewhat higher and more prolonged serum levels of ampicillin and sulbactam can be achieved with the concurrent administration of probenecid.

In patients with impaired renal function the elimination kinetics of ampicillin and sulbactam are similarly affected, hence the ratio of one to the other will remain constant whatever the renal function. The dose of Ampicillin and Sulbactam for Injection in such patients should be administered less frequently in accordance with the usual practice for ampicillin (see DOSAGE AND ADMINISTRATION).

Ampicillin has been found to be approximately 28% reversibly bound to human serum protein and sulbactam approximately 38% reversibly bound.

The following average levels of ampicillin and sulbactam were measured in the tissues and fluids listed:

**TABLE A**  
**Concentration of Ampicillin/Sulbactam in Various Body Tissues and Fluids**

| Fluid or Tissue             | Dose (grams)         | Concentration                             |
|-----------------------------|----------------------|---|
|                             | Ampicillin/Sulbactam | (mcg/mL or mcg/g)<br>Ampicillin/Sulbactam |
| Peritoneal Fluid            | 0.5/0.5 IV           | 7/14                                      |
| Blister Fluid (Cantharides) | 0.5/0.5 IV           | 8/20                                      |
| Tissue Fluid                | 1/0.5 IV             | 8/4                                       |
| Intestinal Mucosa           | 0.5/0.5 IV           | 11/18                                     |
| Appendix                    | 2/1 IV               | 3/40                                      |

Penetration of both ampicillin and sulbactam into cerebrospinal fluid in the presence of inflamed meninges has been demonstrated after IV administration of Ampicillin and Sulbactam for Injection.

The pharmacokinetics of ampicillin and sulbactam in pediatric patients receiving ampicillin and sulbactam are similar to those observed in adults. Immediately after a 15-minute infusion of 50 to 75 mg ampicillin and sulbactam/kg body weight, peak serum and plasma concentrations of 82 to 446 mcg ampicillin/mL and 44 to 203 mcg sulbactam/mL were obtained. Mean half-life values were approximately 1 hour.

### Microbiology

Ampicillin is similar to benzyl penicillin in its bactericidal action against susceptible organisms during the stage of active multiplication. It acts through the inhibition of cell wall mucopeptide biosynthesis. Ampicillin has a broad spectrum of bactericidal activity against many gram-positive and gram-negative aerobic and anaerobic bacteria. (Ampicillin is, however, degraded by beta-lactamases and therefore the spectrum of activity does not normally include organisms which produce these enzymes.)

A wide range of beta-lactamases found in microorganisms resistant to penicillins and cephalosporins have been shown in biochemical studies with cell free bacterial systems to be irreversibly inhibited by sulbactam. Although sulbactam alone possesses little useful antibacterial activity except against the *Neisseriaceae*, whole organism studies have shown that sulbactam restores ampicillin activity against beta-lactamase producing strains. In particular, sulbactam has good inhibitory activity against the clinically important plasmid mediated beta-lactamases most frequently responsible for transferred drug resistance. Sulbactam has no effect on the activity of ampicillin against ampicillin susceptible strains.

The presence of sulbactam in the injection formulation effectively extends the antibiotic spectrum of ampicillin to include many bacteria normally resistant to it and to other beta-lactam antibiotics. Thus, ampicillin and sulbactam possesses the properties of a broad-spectrum antibiotic and a beta-lactamase inhibitor.

While *in vitro* studies have demonstrated the susceptibility of most strains of the following organisms, clinical efficacy for infections other than those included in the indications section has not been documented.

## Gram-Positive Bacteria

*Staphylococcus aureus* (beta-lactamase and non-beta-lactamase producing), *Staphylococcus epidermidis* (beta-lactamase and non-beta-lactamase producing), *Staphylococcus saprophyticus* (beta-lactamase and non-beta-lactamase producing), *Streptococcus faecalis*<sup>1</sup> (*Enterococcus*), *Streptococcus pneumoniae*<sup>1</sup> (formerly *D. pneumoniae*), *Streptococcus pyogenes*<sup>1</sup>, *Streptococcus viridans*<sup>1</sup>.

## Gram-Negative Bacteria

*Hemophilus influenzae* (beta-lactamase and non-beta-lactamase producing), *Moraxella (Branhamella) catarrhalis* (beta-lactamase and non-beta-lactamase producing), *Escherichia coli* (beta-lactamase and non-beta-lactamase producing), *Klebsiella* species (all known strains are beta-lactamase producing), *Proteus mirabilis* (beta-lactamase and non-beta-lactamase producing), *Proteus vulgaris*, *Providencia rettgeri*, *Providencia stuartii*, *Morganella morganii*, and *Neisseria gonorrhoeae* (beta-lactamase and non-beta-lactamase producing).

## Anaerobes

*Clostridium* species<sup>1</sup>, *Peptococcus* species<sup>1</sup>, *Peptostreptococcus* species, *Bacteroides* species, including *B. fragilis*.

<sup>1</sup>These are not beta-lactamase producing strains and, therefore, are susceptible to ampicillin alone.

## Susceptibility Testing

### Diffusion Technique

For the Kirby-Bauer method of susceptibility testing, a 20 mcg (10 mcg ampicillin + 10 mcg sulbactam) diffusion disk should be used.

The method is one outlined in the NCCLS publication M2-A4.<sup>1</sup> With this procedure, a report from the laboratory of "Susceptible" indicates that the infecting organism is likely to respond to ampicillin/sulbactam therapy and a report of "Resistant" indicates that the infecting organism is not likely to respond to therapy. An "Intermediate" susceptibility report suggests that the infecting organism would be susceptible to Ampicillin and Sulbactam for Injection if a higher dosage is used or if the infection is confined to tissues or fluids (e.g., urine) in which high antibiotic levels are attained.

### Dilution Techniques

Broth or agar dilution methods may be used to determine the minimal inhibitory concentration (MIC) value for susceptibility of bacterial isolates to ampicillin/sulbactam. The method used is one outlined in the NCCLS publication M7-A2.<sup>2</sup> Tubes should be inoculated to contain 10<sup>5</sup> to 10<sup>6</sup> organisms/mL or plates "spotted" with 10<sup>4</sup> organisms.

The recommended dilution method employs a constant ampicillin/sulbactam ratio of 2:1 in all tubes with increasing concentrations of ampicillin. MICs are reported in terms of ampicillin concentration in the presence of sulbactam at a constant 2 parts ampicillin to 1 part sulbactam.

### Recommended Ampicillin/Sulbactam, Susceptibility Ranges<sup>\*,†,‡</sup>

|                                   | Resistant | Intermediate | Susceptible |
|-----------------------------------|-----------|--------------|-------------|
| <i>Gram(-) and Staphylococcus</i> | ≤ 11 mm   | 12-13 mm     | ≥ 14 mm     |
| Bauer/Kirby                       | ≥ 32      | 16           | ≤ 8         |
| Zone Sizes                        |           |              |             |
| MIC (mcg of ampicillin/mL)        |           |              |             |
| <i>Hemophilus influenzae</i>      | ≤ 19      | —            | ≥ 20        |
| Bauer/Kirby                       | ≥ 4       | —            | ≤ 2         |
| Zone Sizes                        |           |              |             |
| MIC (mcg of ampicillin/mL)        |           |              |             |

\*The non-beta-lactamase producing organisms which are normally susceptible to ampicillin, such as *Streptococci*, will have similar zone sizes as for ampicillin disks.

†*Staphylococci* resistant to methicillin, oxacillin, or nafcillin must be considered resistant to ampicillin/sulbactam.

‡The quality control cultures should have the following assigned daily ranges for ampicillin/sulbactam:

|                  |              | Disks    | Mode MIC<br>(mcg/mL ampicillin/<br>mcg/mL sulbactam) |
|------------------|--------------|----------|--|
| <i>E. coli</i>   | (ATCC 25922) | 20-24 mm | 2/1  |
| <i>S. aureus</i> | (ATCC 25923) | 29-37 mm | 0.12/0.06  |
| <i>E. coli</i>   | (ATCC 35218) | 13-19 mm | 8/4  |

## INDICATIONS AND USAGE

To reduce the development of drug-resistant bacteria and maintain the effectiveness of Ampicillin and Sulbactam for Injection and other antibacterial drugs, Ampicillin and Sulbactam for Injection should be used only to treat or prevent infections that are proven or strongly suspected to be caused by susceptible bacteria. When culture and susceptibility information are available, they should be considered in selecting or modifying antibacterial therapy. In the absence of such data, local epidemiology and susceptibility patterns may contribute to the empiric selection of therapy.

Ampicillin and Sulbactam for Injection is indicated for the treatment of infections due to susceptible strains of the designated microorganisms in the conditions listed below.

**Skin and Skin Structure Infections** caused by beta-lactamase producing strains of *Staphylococcus aureus*, *Escherichia coli*,<sup>2</sup> *Klebsiella* spp.<sup>2</sup> (including *K. pneumoniae*<sup>2</sup>), *Proteus mirabilis*,<sup>2</sup> *Bacteroides fragilis*,<sup>2</sup> *Enterobacter* spp.,<sup>2</sup> and *Acinetobacter calcoaceticus*<sup>2</sup>.

NOTE: For information on use in pediatric patients see PRECAUTIONS—Pediatric Use and CLINICAL STUDIES sections.

**Intra-Abdominal Infections** caused by beta-lactamase producing strains of *Escherichia coli*, *Klebsiella* spp. (including *K. pneumoniae*<sup>2</sup>), *Bacteroides* spp. (including *B. fragilis*), and *Enterobacter* spp.<sup>2</sup>

**Gynecological Infections** caused by beta-lactamase producing strains of *Escherichia coli*<sup>2</sup> and *Bacteroides* spp.<sup>2</sup> (including *B. fragilis*<sup>2</sup>).

While Ampicillin and Sulbactam for Injection is indicated only for the conditions listed above, infections caused by ampicillin-susceptible organisms are also amenable to treatment with Ampicillin and Sulbactam for Injection due to its ampicillin content. Therefore, mixed infections caused by ampicillin-susceptible organisms and beta-lactamase producing organisms susceptible to Ampicillin and Sulbactam for Injection should not require the addition of another antibiotic.

Appropriate culture and susceptibility tests should be performed before treatment in order to isolate and identify the organisms causing infection and to determine their susceptibility to Ampicillin and Sulbactam for Injection.

Therapy may be instituted prior to obtaining the results from bacteriological and susceptibility studies, when there is reason to believe the infection may involve any of the beta-lactamase producing organisms listed above in the indicated organ systems. Once the results are known, therapy should be adjusted if appropriate.

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<sup>2</sup>Efficacy for this organism in this organ system was studied in fewer than 10 infections.

## CONTRAINDICATIONS

The use of Ampicillin and Sulbactam for Injection is contraindicated in individuals with a history of hypersensitivity reactions to any of the penicillins.

## WARNINGS

SERIOUS AND OCCASIONALLY FATAL HYPERSENSITIVITY (ANAPHYLACTIC) REACTIONS HAVE BEEN REPORTED IN PATIENTS ON PENICILLIN THERAPY. THESE REACTIONS ARE MORE APT TO OCCUR IN INDIVIDUALS WITH A HISTORY OF PENICILLIN HYPERSENSITIVITY AND/OR HYPERSENSITIVITY REACTIONS TO MULTIPLE ALLERGENS. THERE HAVE BEEN REPORTS OF INDIVIDUALS WITH A HISTORY OF PENICILLIN HYPERSENSITIVITY WHO HAVE EXPERIENCED SEVERE REACTIONS WHEN TREATED WITH CEPHALOSPORINS. BEFORE THERAPY WITH A PENICILLIN, CAREFUL INQUIRY SHOULD BE MADE CONCERNING PREVIOUS HYPERSENSITIVITY REACTIONS TO PENICILLINS, CEPHALOSPORINS, AND OTHER ALLERGENS. IF AN ALLERGIC REACTION OCCURS, AMPICILLIN AND SULBACTAM FOR INJECTION SHOULD BE DISCONTINUED AND THE APPROPRIATE THERAPY INSTITUTED. SERIOUS ANAPHYLACTOID REACTIONS REQUIRE IMMEDIATE EMERGENCY TREATMENT WITH EPINEPHRINE. OXYGEN, INTRAVENOUS STEROIDS, AND AIRWAY MANAGEMENT, INCLUDING INTUBATION, SHOULD ALSO BE ADMINISTERED AS INDICATED.

**Pseudomembranous colitis has been reported with nearly all antibacterial agents, including Ampicillin and Sulbactam for Injection, and has ranged in severity from mild to life-threatening. Therefore, it is important to consider this diagnosis in patients who present with diarrhea subsequent to the administration of antibacterial agents.**

Treatment with antibacterial agents alters the normal flora of the colon and may permit overgrowth of clostridia. Studies indicate that toxin produced by *Clostridium difficile* is one primary cause of "antibiotic-associated colitis."

Mild cases of pseudomembranous colitis usually respond to drug discontinuation alone. In moderate to severe cases, consideration should be given to management with fluids and electrolytes, protein supplementation and treatment with an antibacterial drug clinically effective against *C. difficile* colitis.

## PRECAUTIONS

### General

Prescribing Ampicillin and Sulbactam for Injection in the absence of a proven or strongly suspected bacterial infection or a prophylactic indication is unlikely to provide benefit to the patient and increases the risk of the development of drug-resistant bacteria.

A high percentage of patients with mononucleosis who receive ampicillin develop a skin rash. Thus, ampicillin class antibiotics should not be administered to patients with mononucleosis. In patients treated with ampicillin and sulbactam the possibility of superinfections with mycotic or bacterial pathogens should be kept in mind during therapy. If superinfections occur (usually involving *Pseudomonas* or *Candida*), the drug should be discontinued and/or appropriate therapy instituted.

### **Information for Patients**

Patients should be counseled that antibacterial drugs including Ampicillin and Sulbactam for Injection should only be used to treat bacterial infections. They do not treat viral infections (e.g., the common cold). When Ampicillin and Sulbactam for Injection is prescribed to treat a bacterial infection, patients should be told that although it is common to feel better early in the course of therapy, the medication should be taken exactly as directed. Skipping doses or not completing the full course of therapy may (1) decrease the effectiveness of the immediate treatment and (2) increase the likelihood that bacteria will develop resistance and will not be treatable by Ampicillin and Sulbactam for Injection or other antibacterial drugs in the future.

### **Drug Interactions**

Probenecid decreases the renal tubular secretion of ampicillin and sulbactam. Concurrent use of probenecid with ampicillin and sulbactam may result in increased and prolonged blood levels of ampicillin and sulbactam. The concurrent administration of allopurinol and ampicillin increases substantially the incidence of rashes in patients receiving both drugs as compared to patients receiving ampicillin alone. It is not known whether this potentiation of ampicillin rashes is due to allopurinol or the hyperuricemia present in these patients. There are no data with ampicillin and sulbactam and allopurinol administered concurrently. Ampicillin and sulbactam and aminoglycosides should not be reconstituted together due to the *in vitro* inactivation of aminoglycosides by the ampicillin component of Ampicillin and Sulbactam for Injection.

### **Drug/Laboratory Test Interactions**

Administration of ampicillin and sulbactam will result in high urine concentration of ampicillin. High urine concentrations of ampicillin may result in false positive reactions when testing for the presence of glucose in urine using Clinitest™, Benedict's Solution or Fehling's Solution. It is recommended that glucose tests based on enzymatic glucose oxidase reactions (such as Clinistix™) be used.

Following administration of ampicillin to pregnant women, a transient decrease in plasma concentration of total conjugated estriol, estriol-glucuronide, conjugated estrone and estradiol has been noted. This effect may also occur with ampicillin and sulbactam.

### **Carcinogenesis and Mutagenesis and Impairment of Fertility**

Long-term studies in animals have not been performed to evaluate carcinogenic or mutagenic potential.

### **Pregnancy**

#### **Pregnancy Category B**

Reproduction studies have been performed in mice, rats, and rabbits at doses up to ten (10) times the human dose and have revealed no evidence of impaired fertility or harm to the fetus due to Ampicillin and Sulbactam for Injection. There are, however, no adequate and well controlled studies in pregnant women. Because animal reproduction studies are not always predictive of human response, this drug should be used during pregnancy only if clearly needed. (See Drug/Laboratory Test Interactions.)

### **Labor and Delivery**

Studies in guinea pigs have shown that intravenous administration of ampicillin decreased the uterine tone, frequency of contractions, height of contractions, and duration of contractions. However, it is not known whether the use of Ampicillin and Sulbactam for Injection in humans during labor or delivery has immediate or delayed adverse effects on the fetus, prolongs the duration of labor, or increases the likelihood that forceps delivery or other obstetrical intervention or resuscitation of the newborn will be necessary.

### **Nursing Mothers**

Low concentrations of ampicillin and sulbactam are excreted in the milk; therefore, caution should be exercised when Ampicillin and Sulbactam for Injection is administered to a nursing woman.

### **Pediatric Use**

The safety and effectiveness of Ampicillin and Sulbactam for Injection have been established for pediatric patients one year of age and older for skin and skin structure infections as approved in adults. Use of Ampicillin and Sulbactam for Injection in pediatric patients is supported by evidence from adequate and well-controlled studies in adults with additional data from pediatric pharmacokinetic studies, a controlled clinical trial conducted in pediatric patients and post-marketing adverse events surveillance. (See CLINICAL PHARMACOLOGY, INDICATIONS AND USAGE, ADVERSE REACTIONS, DOSAGE AND ADMINISTRATION, and CLINICAL STUDIES sections.)

The safety and effectiveness of Ampicillin and Sulbactam for Injection have not been established for pediatric patients for intra-abdominal infections.

## **ADVERSE REACTIONS**

### **Adult Patients**

Ampicillin and Sulbactam for Injection is generally well tolerated. The following adverse reactions have been reported.

#### **Local Adverse Reactions**

Pain at IM injection site—16%

Pain at IV injection site—3%

Thrombophlebitis—3%

#### **Systemic Adverse Reactions**

The most frequently reported adverse reactions were diarrhea in 3% of the patients and rash in less than 2% of the patients.

Additional systemic reactions reported in less than 1% of the patients were: itching, nausea, vomiting, candidiasis, fatigue, malaise, headache, chest pain, flatulence, abdominal distension, glossitis, urine retention, dysuria, edema, facial swelling, erythema, chills, tightness in throat, substernal pain, epistaxis and mucosal bleeding.

### **Pediatric Patients**

Available safety data for pediatric patients treated with Ampicillin and Sulbactam for Injection demonstrate a similar adverse events profile to those observed in adult patients. Additionally, atypical lymphocytosis has been observed in one pediatric patient receiving Ampicillin and Sulbactam for Injection.

### **Adverse Laboratory Changes**

Adverse laboratory changes without regard to drug relationship that were reported during clinical trials were:

#### **Hepatic**

Increased AST (SGOT), ALT (SGPT), alkaline phosphatase, and LDH.

#### **Hematologic**

Decreased hemoglobin, hematocrit, RBC, WBC, neutrophils, lymphocytes, platelets and increased lymphocytes, monocytes, basophils, eosinophils, and platelets.

#### **Blood Chemistry**

Decreased serum albumin and total proteins.

#### **Renal**

Increased BUN and creatinine.

#### **Urinalysis**

Presence of RBCs and hyaline casts in urine.

The following adverse reactions have been reported with ampicillin-class antibiotics and can also occur with Ampicillin and Sulbactam for Injection.

### **Gastrointestinal**

Gastritis, stomatitis, black "hairy" tongue and enterocolitis. Onset of pseudomembranous colitis symptoms may occur during or after antibiotic treatment. (See WARNINGS.)

### **Hypersensitivity Reactions**

Urticaria, erythema multiforme, and an occasional case of exfoliative dermatitis have been reported. These reactions may be controlled with antihistamines and, if necessary, systemic corticosteroids. Whenever such reactions occur, the drug should be discontinued, unless the opinion of the physician dictates otherwise. Serious and occasional fatal hypersensitivity (anaphylactic) reactions can occur with a penicillin. (See WARNINGS.)

### **Hematologic**

In addition to the adverse laboratory changes listed above for Ampicillin and Sulbactam for Injection, agranulocytosis has been reported during therapy with penicillins. All of these reactions are usually reversible on discontinuation of therapy and are believed to be hypersensitivity phenomena. Some individuals have developed positive direct Coombs Tests during treatment with Ampicillin and Sulbactam for Injection, as with other beta-lactam antibiotics.

## OVERDOSAGE

Neurological adverse reactions, including convulsions, may occur with the attainment of high CSF levels of beta-lactams. Ampicillin may be removed from circulation by hemodialysis. The molecular weight, degree of protein binding and pharmacokinetics profile of sulbactam suggest that this compound may also be removed by hemodialysis.

## CLINICAL STUDIES

### Skin and Skin Structure Infections in Pediatric Patients

Data from a controlled clinical trial conducted in pediatric patients provided evidence supporting the safety and efficacy of Ampicillin and Sulbactam for Injection for the treatment of skin and skin structure infections. Of 99 pediatric patients evaluable for clinical efficacy, 60 patients received a regimen containing intravenous Ampicillin and Sulbactam for Injection, and 39 patients received a regimen containing intravenous cefuroxime. This trial demonstrated similar outcomes (assessed at an appropriate interval after discontinuation of all antimicrobial therapy) for ampicillin/sulbactam- and cefuroxime-treated patients:

| <u>Therapeutic Regimen</u> | <u>Clinical Success</u> | <u>Clinical Failure</u> |
|----------------------------|-------------------------|-------------------------|
| Ampicillin/Sulbactam       | 51/60 (85%)             | 9/60 (15%)              |
| Cefuroxime                 | 34/39 (87%)             | 5/39 (13%)              |

Most patients received a course of oral antimicrobials following initial treatment with intravenous administration of parenteral antimicrobials. The study protocol required that the following three criteria be met prior to transition from intravenous to oral antimicrobial therapy: 1) receipt of a minimum of 72 hours of intravenous therapy; 2) no documented fever for prior 24 hours; and 3) improvement or resolution of the signs and symptoms of infection.

The choice of oral antimicrobial agent used in this trial was determined by susceptibility testing of the original pathogen, if isolated, to oral agents available. The course of oral antimicrobial therapy should not routinely exceed 14 days.

## DOSAGE AND ADMINISTRATION

Ampicillin and Sulbactam for Injection may be administered by either the IV or deep IM routes (**following dilution**). **HOWEVER, THE INTENT OF THE PHARMACY BULK PACKAGE IS FOR PREPARATION OF SOLUTIONS FOR IV INFUSION ONLY.**

For IV administration, the dose can be given by slow intravenous injection over at least 10-15 minutes or can also be delivered in greater dilutions with 50-100 mL of a compatible diluent as an intravenous infusion over 15-30 minutes.

The recommended adult dosage of ampicillin and sulbactam is 1.5 g (1 g ampicillin as the sodium salt plus 0.5 g sulbactam as the sodium salt) to 3 g (2 g ampicillin as the sodium salt plus 1 g sulbactam as the sodium salt) every six hours. This 1.5 to 3 g range represents the total of ampicillin content plus the sulbactam content of Ampicillin and Sulbactam for Injection, and corresponds to a range of 1 g ampicillin/0.5 g sulbactam to 2 g ampicillin/1 g sulbactam. The total dose of sulbactam should not exceed 4 grams per day.

### Pediatric Patients 1 Year of Age or Older

The recommended daily dose of Ampicillin and Sulbactam for Injection in pediatric patients is 300 mg per kg of body weight administered via intravenous infusion in equally divided doses every 6 hours. This 300 mg/kg/day dosage represents the total ampicillin content plus the sulbactam content of Ampicillin and Sulbactam for Injection, and corresponds to 200 mg ampicillin/100 mg sulbactam per kg per day. The safety and efficacy of Ampicillin and Sulbactam for Injection administered via intramuscular injection in pediatric patients have not been established.

Pediatric patients weighing 40 kg or more should be dosed according to adult recommendations, and the total dose of sulbactam should not exceed 4 grams per day. The course of intravenous therapy should not routinely exceed 14 days. In clinical trials, most children received a course of oral antimicrobials following initial treatment with intravenous Ampicillin and Sulbactam for Injection. (See CLINICAL STUDIES section.)

### Impaired Renal Function

In patients with impairment of renal function the elimination kinetics of ampicillin and sulbactam are similarly affected, hence the ratio of one to the other will remain constant whatever the renal function. The dose of Ampicillin and Sulbactam for Injection in such patients should be administered less frequently in accordance with the usual practice for ampicillin and according to the following recommendations:

Ampicillin and Sulbactam for Injection Dosage Guide For Patients With Renal Impairment

| <b>Creatinine Clearance</b><br><b>(mL/min/1.73m<sup>2</sup>)</b> | <b>Ampicillin/Sulbactam Half-Life (Hours)</b> | <b>Recommended</b><br><b>Ampicillin/Sulbactam Dosage</b> |
|--|---|--|
| ≥30  | 1   | 1.5-3 g q 6h-q 8h  |
| 15-29  | 5   | 1.5-3 g q 12h  |
| 5-14   | 9   | 1.5-3 g q 24h  |

When only serum creatinine is available, the following formula (based on sex, weight, and age of the patient) may be used to convert this value into creatinine clearance. The serum creatinine should represent a steady state of renal function.

|         |  |
|---------|--|
| Males   | $\frac{\text{weight (kg)} \times (140 - \text{age})}{72 \times \text{serum creatinine}}$ |
| Females | 0.85 x above value   |

### Compatibility, Reconstitution and Stability

When concomitant therapy with aminoglycosides is indicated, ampicillin and sulbactam and aminoglycosides should be reconstituted and administered separately, due to the *in vitro* inactivation of aminoglycosides by any of the aminopenicillins.

### Directions for Use

#### General Dissolution Procedures

Ampicillin and Sulbactam for Injection sterile powder for intravenous use may be reconstituted with any of the compatible diluents described in this insert. Solutions should be allowed to stand after dissolution to allow any foaming to dissipate in order to permit visual inspection for complete solubilization.

#### Preparation for Intravenous Use

Ampicillin and Sulbactam for Injection concentrations between 3 and 45 mg ampicillin and sulbactam (2 to 30 mg ampicillin/1 to 15 mg sulbactam/mL) are recommended for intravenous use.

#### Directions for Proper Use of Pharmacy Bulk Package

The 15 gram vial may be reconstituted with either 92 mL Sterile Water for Injection or 0.9% Sodium Chloride Injection. The diluent should be added in two separate aliquots in a suitable work area, such as a laminar flow hood. Add 50 mL of solution, shake to dissolve. Then add an additional 42 mL and shake. The solution should be allowed to stand after dissolution to allow any foaming to dissipate in order to permit visual inspection for complete solubilization. The resultant solution will have a final concentration of approximately 100 mg/mL ampicillin and 50 mg/mL sulbactam. The closure may be penetrated only one time after reconstitution, if needed, using a suitable sterile transfer device or dispensing set that allows for measured dispensing of the contents.

After reconstitution, use within two hours if stored at room temperature, or within four hours if stored under refrigeration.

|   |
|---|
| <b>Reconstituted Bulk Solution Should Not be Used For Direct Infusion</b> |
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If the reconstituted bulk solution is stored for less than one hour at room temperature (20°C/68°F) prior to further dilution, the use periods indicated in Table A apply for the diluted solutions.

If the bulk solution is stored for one to two hours at room temperature (20°C/68°F) and then diluted with Sterile Water for Injection or 0.9% Sodium Chloride Injection to the following concentrations, the use periods indicated in Table B apply.

Any unused portions of solution that remain after the indicated time periods should be discarded.

**TABLE A**  
**Maximum Concentration (mg/mL)**

| Diluent                        | Ampicillin/Sulbactam | Use Periods  |
|--------------------------------|----------------------|--------------|
| Sterile Water for Injection    | 45 (30/15)           | 8 hrs @ 21°C |
|                                | 45 (30/15)           | 48 hrs @ 4°C |
|                                | 30 (20/10)           | 72 hrs @ 4°C |
| 0.9% Sodium Chloride Injection | 45 (30/15)           | 8 hrs @ 21°C |
|                                | 45 (30/15)           | 48 hrs @ 4°C |
|                                | 30 (20/10)           | 72 hrs @ 4°C |
| 5% Dextrose Injection          | 30 (20/10)           | 2 hrs @ 21°C |
|                                | 30 (20/10)           | 4 hrs @ 4°C  |
|                                | 3 (2/1)              | 2 hrs @ 21°C |
| Lactated Ringer's Injection    | 45 (30/15)           | 8 hrs @ 21°C |
|                                | 45 (30/15)           | 24 hrs @ 4°C |
| M/6 Sodium Lactate Injection   | 45 (30/15)           | 8 hrs @ 21°C |
|                                | 45 (30/15)           | 12 hrs @ 4°C |
| 5% Dextrose in 0.45% Saline    | 3 (2/1)              | 4 hrs @ 21°C |



|                  |            |              |
|------------------|------------|--------------|
|                  | 15 (10/5)  | 4 hrs @ 4°C  |
| 10% Invert Sugar | 3 (2/1)    | 4 hrs @ 21°C |
|                  | 30 (20/10) | 3 hrs @ 4°C  |

**TABLE B**  
**Maximum Concentration (mg/mL)**

| <b>IV Solution</b>                  | <b>Ampicillin/Sulbactam</b> | <b>Use Periods</b> |
|-------------------------------------|-----------------------------|--------------------|
| Sterile Water for Injection, USP    | 45 (30/15)                  | 4 hrs @ 21°C       |
|                                     | 45 (30/15)                  | 24 hrs @ 4°C       |
| 0.9% Sodium Chloride Injection, USP | 45 (30/15)                  | 4 hrs @ 21°C       |
|                                     | 45 (30/15)                  | 24 hrs @ 4°C       |

## HOW SUPPLIED

Ampicillin and Sulbactam for Injection, USP, a sterile off-white dry powder, is available in a 15 gram Pharmacy Bulk Package. Ampicillin and Sulbactam for Injection, USP containing ampicillin sodium equivalent to 10 grams of ampicillin and sulbactam sodium equivalent to 5 grams of sulbactam—packaged individually (NDC 10019-632-03).

## Storage

**Prior to reconstitution store dry powder at 20°-25°C (68°-77°F) [see USP Controlled Room Temperature].**

## ANIMAL PHARMACOLOGY

While reversible glycogenesis was observed in laboratory animals, this phenomenon was dose- and time-dependent and is not expected to develop at the therapeutic doses and corresponding plasma levels attained during the relatively short periods of combined ampicillin/sulbactam therapy in man.

## REFERENCES

1. National Committee for Clinical Laboratory Standards, *Performance Standards for Antimicrobial Disk Susceptibility Tests*-Fourth Edition. Approved Standard NCCLS Document M2-A4, Vol. 10, No. 7 NCCLS, Villanova, PA, April 1990.
2. National Committee for Clinical Laboratory Standards, *Methods for Dilution Antimicrobial Susceptibility Tests for Bacteria that Grow Aerobically*, Second Edition. Approved Standard NCCLS Document M7-A2, Vol. 10, No. 8 NCCLS, Villanova, PA, April 1990.

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